Docket No. 1829-4004US1

<u>IN THE UNITED STATES PATENT AND TRADEMARK OFFICE</u>

cant(s):

SUSAN A. SLAUGENHAUPT, ET AL.

Group Art Unit:

1645

Serial No .:

10/041,856

Examiner:

Not Yet Assigned

Filed:

January 7, 2002

For:

GENE FOR IDENTIFYING INDIVIDUALS WITH FAMILIAL DYSAUTONOMIA

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INFORMATION DISCLOSURE STATEMENT

Pursuant to Rule 56, applicant hereby calls the attention of the Patent Office to the references

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By:

Dated: July 17, 2003

Respectfully submitted,
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INFORM TION DISCLOSURE CITATION Applicant: Susan Shaugenhaup, et al. Filing Date: Group Art Unit: January 7, 2002 TBA	OIPE	FORM PTO)-1449		Attorney Docket: 1829-4004US1	Serial N 10/041,		*
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U.S. PATENT DOCUMENTS U.S. PATENT DOCUMENTS U.S. PATENT DOCUMENTS Patent Number Date Name Class Sub-Class Fill 5,387,506 02/07/95 A. Blumenfeld, et al. 435 6 5 FOREIGN PATENT DOCUMENTS Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 D.	INFORM	TION DISCLO	OSURE CITA	ATION			Art I Init:	
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FOREIGN PATENT DOCUMENTS Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 COTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								T1111 TO 1
FOREIGN PATENT DOCUMENTS Examiner Patent Number Date Country Class Sub-Class Tr.	Initial			A 701				Filing Date
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL "Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		5,387,506	02/07/95	A. Blum	enicia, et al.	435	66	5/29/92
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT CO7 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL "Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								
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Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT CO7 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.	MILE							
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL "Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,						
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT CO7 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes In Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								
Examiner Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT CO7 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associ protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								
Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.			F	OREIGN I	PATENT DOCUMENTS			
Initial Patent Number Date Country Class Sub-Class Tr. WO 99/25730 5/27/99 PCT C07 K 7/06 EP 1 225 232 A2 7/24/02 Europe C12 Q 1/68 OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the 1 B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.	Examiner		Publication					
EP 1 225 232 A2 7/24/02 Europe C12 Q 1 / 68		Patent Number						Translatio
OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes F Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.				*				N/A
OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes F. Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		EP 1 225 232 A2	7/24/02		Europe	C12 Q	1 / 68	N/A
OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Papers, etc.) L. Cohen, et al., "IKAP is a scaffold protein of the I B kinase complex," Nature, 395:292-296, 17 Septem S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes I Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-associprotein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								☐ Yes ☐ N
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S. A. Slaugenhaupt, et al., "Tissue-Specific Express of a Splicing Mutation in the IKBKAP Gene Causes Education Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-association of the IKBKAP) mRNA," retrieved from http://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		7 61	WIZAD '	CC. 1.1	:	205	202 206 17 9	
Dysautonomia," Am. J. Hum. Genet., 68:598-605, 2001. S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-association of the IKBKAP) mRNA," retrieved from http://www.EBI.AC.UK, Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.								
S. A. Slaugenhaupt, et al., Database EMBL 'Online! EBI; "Homo sapiens IkappaBkinase complex-association (IKBKAP) mRNA," retrieved from http://www.EBI.AC.UK , Database accession no.: AF15341 January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," <a "familial="" 2001="" 2001.="" 2001.<="" 28,="" 68:="" 753-758,="" accession="" af15341="" al.,="" am.="" anderson,="" by="" caused="" database="" dysautonomia="" et="" february="" from="" gene,"="" href="https://www.ebi.ncbi.ncbi.ncbi.ncbi.ncbi.ncbi.ncbi.nc</td><td></td><td></td><td></td><td></td><td></td><td>utation in the IKE</td><td>BKAP Gene Cau</td><td>uses Familial</td></tr><tr><td>protein (IKBKAP) mRNA," http:="" hum.="" ikap="" is="" j.="" january="" l.="" mutations="" no.:="" of="" retrieved="" s.="" td="" the="" updated="" www.ebi.ac.uk,=""><td></td><td>Dysautonomia,"</td><td>Am. J. Hum. C</td><td><u>ienet.</u>, 68::</td><td>598-605, 2001.</td><td></td><td></td><td></td>		Dysautonomia,"	Am. J. Hum. C	<u>ienet.</u> , 68::	598-605, 2001.			
January 2001 updated February 28, 2001. S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		S. A. Slaugenha	upt, et al., Data	base EMB	L 'Online! EBI; "Homo sa	piens IkappaBkir	nase complex-a	ssociated
S. L. Anderson, et al., "Familial Dysautonomia is Caused by Mutations of the IKAP Gene," Am. J. Hum. 68: 753-758, 2001.		protein (IKBKAP) mRNA," retrieved from HTTP://www.EBI.AC.UK, Database accession no.: AF1534						53419, 2
68: 753-758, 2001.								
							Gene," <u>Am. J. I</u>	Ium. Genet.,
Examiner Date Considered		68: 753-758, 20	01.					
Examiner Date Considered								
Examiner Date Considered								
Examiner Date Considered								P
Date Considered	Examiner				Date Considered	i		
EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP §609.								

PATENT COOPERATION TREATY



PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	(Form PCT/ISA/220) as well as, where applicable, item 5 below.		
1829-4004PC International application No.	International filing date (day/month/year)	/ (Endingt) Princip Data (day/orgath/agail	
	international filing date (day/month/year)	(Earliest) Priority Date (day/month/year)	
PCT/US 02/00473	06/01/2001		
DYSAUTONOMIA FOUNDATION I	NC.		
DISAUTUNUMIA FOUNDATION I	NC.		
This International Search Report has been according to Article 18. A copy is being tra	n prepared by this International Searching Auth ansmitted to the International Bureau.	nority and is transmitted to the applicant	
This International Search Report consists [X] It is also accompanied by	of a total of6 sheets. a copy of each prior art document cited in this	report.	
Basis of the report	1-A		
language in which it was filed, unl	international search was carried out on the bases otherwise indicated under this item.	sis of the international application in the	
the international search w Authority (Rule 23.1(b)).	ras carried out on the basis of a translation of the	he international application furnished to this	
was carried out on the basis of the	e sequence listing:	ternational application, the international search	
	onal application in written form.	_	
t-mand	ernational application in computer readable form	n. ·	
	this Authority in written form.		
	this Authority in computer readble form.		
international application a	osequently furnished written sequence listing d is filed has been furnished.	oes not go beyond the disclosure in the	
X the statement that the info	ormation recorded in computer readable form is	s identical to the written sequence listing has been	
2. X Certain claims were fou	nd unsearchable (See Box I).		
3. Unity of invention is lack	king (see Box II).		
4. With regard to the title,			
X the text is approved as su	bmitted by the applicant.		
the text has been establis	hed by this Authority to read as follows:		
5. With regard to the abstract,			
	.h.milled by the englished		
the text is approved as su the text has been establis within one month from the	brifficed by the applicant. hed, according to Rule 38.2(b), by this Authori date of mailing of this international search rep	ty as it appears in Box III. The applicant may, oort, submit comments to this Authority.	
6. The figure of the drawings to be publ	ished with the abstract is Figure No.		
as suggested by the appli	cant.	X None of the figures.	
because the applicant fail	ed to suggest a figur .		
because this figure better	characterizes the invention.		

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 02/00473

a. CLASSIFICATION OF SUBJECT MATTER IPC 7 C12Q1/68 C07K14/47 C12N15/63 C12N15/10 A01K67/027

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, EMBL, MEDLINE, EMBASE, PAJ, BIOSIS, WPI Data

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
K	COHEN L HENZEL W J BAEUERLE P A: "IKAP is a scaffold protein of the IkappaB kinase complex" NATURE, MACMILLAN JOURNALS LTD. LONDON, GB, vol. 395, 17 September 1998 (1998-09-17), pages 292-296, XP002956454 ISSN: 0028-0836 cited in the application page 296, left-hand column, paragraph 3 -right-hand column, paragraph 3 figure 2	1,2,6,8, 10-13, 29,30,43
	-/	

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"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or	 "T" tater document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	 "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search	Date of mailing of the international search report
2 June 2003	24/06/2003
Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Ulbrecht, M

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INTERNATIONAL'SEARCH REPORT

International Application No
PCT/US 02/00473

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	Citation of decument with indication where constraints of the relevant	
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	W0 99 25730 A (TULARIK INC) 27 May 1999 (1999-05-27) page 8, line 6 -page 9, line 29 claims 1-5	1,2,6,8, 10-13, 29,30,43
X	SEQ ID Nos. 1 and 2 DATABASE EMBL 'Online! EBI; 2 January 2001 (2001-01-02) SLAUGENHAUPT S.A. ET AL.: "Homo sapiens	1,2,6,8, 10-13, 29,30,43
	IkappaBkinase complex-associated protein (IKBKAP) mRNA" retrieved from HTTP://WWW.EBI.AC.UK Database accession no. AF153419 XP002240942 the whole document	
Ρ,Χ	SLAUGENHAUPT SUSAN A ET AL: "Tissue-specific expression of a splicing mutation in the IKBKAP gene causes familial dysautonomia" AMERICAN JOURNAL OF HUMAN GENETICS, AMERICAN SOCIETY OF HUMAN GENETICS, CHICAGO, IL, US, vol. 68, no. 3, March 2001 (2001-03), pages 598-605, XP002188142 ISSN: 0002-9297 the whole document	1-43
Ρ,Χ	ANDERSON S L ET AL: "FAMILIAL DYSAUTONOMIA IS CAUSED BY MUTATIONS OF THE IKAP GENE" AMERICAN JOURNAL OF HUMAN GENETICS, AMERICAN SOCIETY OF HUMAN GENETICS, CHICAGO, IL, US, vol. 68, no. 3, March 2001 (2001-03), pages 753-758, XP009005056 ISSN: 0002-9297 the whole document	1-43
E	EP 1 225 232 A (ANDERSON SILVIA L ; RUBIN BERISH Y (US)) 24 July 2002 (2002-07-24)	1-9,15, 20, 28-30, 42,43
A	the whole document US 5 387 506 A (BLUMENFELD ANAT ET AL)	1-43
n	7 February 1995 (1995-02-07) the whole document	1-43

INTERNATIONAL'SEARCH REPORT

Information on patent family members

International Application No PCT/US 02/00473

	atent document I in search report		Publication date		Patent family member(s)	Publication date
WO	9925730	A	27-05-1999	US AU WO US	5891719 A 1525499 A 9925730 A1 6172195 B1	06-04-1999 07-06-1999 27-05-1999 09-01-2001
EP	1225232	Α	24-07-2002	CA EP US	2366046 A1 1225232 A2 2002168656 A1	17-07-2002 24-07-2002 14-11-2002
US	5387506	A	07-02-1995	AU CA GB IL WO US US US	4390293 A 2136859 A1 2282446 A ,B 105818 A 120076 A 9324657 A2 6262250 B1 5998133 A 2002025528 A1 9303670 A	30-12-1993 09-12-1993 05-04-1995 04-01-1998 28-10-1999 09-12-1993 17-07-2001 07-12-1999 28-02-2002 01-08-1994

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 34,35

Present claims 34 and 35 relate to a product defined by reference to a desirable characteristic or property, namely oligonucleotides which hybridise to one opr more additional mutant or wild-type genes associated with additional genetic diseases. The claims cover all products having this characteristic or property, whereas the application provides neither support within the meaning of Art. 6 PCT nordisclosure within the meaning of Article 5 PCT for any such product. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search is impossible. Independent of the above reasoning, the claims also lack clarity (Art. 6 PCT). An attempt is made to define the product by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the subject-matter of claims 34 and 35 has not been searched at all.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

International application No. PCT/US 02/00473

Box I Observati ns wh re certain claims were found unsearchable (Continuation of item 1 of first sh et)
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
2. X Claims Nos.: 34,35 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically: see FURTHER INFORMATION sheet PCT/ISA/210
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This International Searching Authority found multiple inventions in this international application, as follows:
As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest The additional search fees were accompanied by the applicant's protest. No protest accompanied the payment of additional search fees.